



Food and Drug Administration Rockville, MD 20857

February 22, 2005

FILE COPY

Robert W. Pollock, Vice President Lachman Consultant Services, Inc. 1600 Stewart Avenue Westbury, NY 11590

Dear Mr. Pollock:

Your petition requesting the Food and Drug Administration to make a determination that the drug product, Prednisolone Sodium Phosphate Solution, in the strengths of 10 mg/5 mL, 20 mg/5 ml and 25 mg/5 mL (equivalent prednisolone base), was received by this office on 02/22/2005. It was assigned docket number 2005P-0080/CP 1 and it was filed on 02/22/2005. Please refer to this docket number in future correspondence on this subject with the Agency.

Please note that the acceptance of the petition for filing is a procedural matter in that it in no way reflects an agency decision on the substantive merits of the petition.

Sincerely,

Ľyle D. Jáffe

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Office of Management

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